

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization International Bureau



(43) International Publication Date
18 March 2004 (18.03.2004)

PCT

(10) International Publication Number
WO 2004/022096 A1

(51) International Patent Classification⁷: **A61K 39/395, C07K 16/36**

(21) International Application Number:
PCT/US2003/027808

(22) International Filing Date:
5 September 2003 (05.09.2003)

(25) Filing Language: **English**

(26) Publication Language: **English**

(30) Priority Data:
60/408,571 6 September 2002 (06.09.2002) US
60/469,189 9 May 2003 (09.05.2003) US

(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

(71) Applicant (for all designated States except US): **ALEXION PHARMACEUTICALS, INC. [US/US]; 352 Knotter Drive, Cheshire, CT 06410 (US)**

(72) Inventor; and

(75) Inventor/Applicant (for US only): **WANG, Yi [US/US]; 270 Sylvan Valley Road, Orange, CT 06477 (US)**

(74) Agent: **FARBER, Mark; Alexion Pharmaceuticals, Inc., 352 Knotter Drive, Cheshire, CT 06410 (US)**

Published:

- with international search report
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

WO 2004/022096 A1

(54) Title: **METHOD OF TREATMENT OF ASTHMA USING ANTIBODIES TO COMPLEMENT COMPONENT C5**

(57) Abstract: A method of treatment of asthma with a compound which binds to or otherwise blocks the generation and/or activity of one or more complement components such as, for example, a complement-inhibiting antibody. The compound can be administered prophylactically to prevent asthma attacks in known asthmatic individuals or as a therapeutic regimen during asthma attacks.

5 METHOD OF TREATMENT OF ASTHMA USING ANTIBODIES TO COMPLEMENT COMPONENT C5

BACKGROUND

TECHNICAL FIELD

This disclosure relates to a method of treating asthma using a compound
10 which binds to or otherwise blocks the generation and/or activity of one or more
complement components such as, for example, a complement-inhibiting
antibody.

DESCRIPTION OF THE RELATED ART

Asthma, bronchitis and emphysema are known collectively as Chronic Obstructive Pulmonary Diseases. These diseases are characterized as generalized airways obstruction, especially of small airways, associated with varying degrees of symptoms of chronic bronchitis, asthma, and emphysema. These diseases may often coexist in an individual, and it may be difficult to determine the primary cause of an airway obstruction. Airway obstruction is defined as an increased resistance to airflow during forced expiration. Obstruction of large airways may also occur in these diseases, particularly in asthma.

Asthma is a reversible obstructive pulmonary disorder caused by an airway hyper-responsiveness to specific and/ or non-specific stimuli. Asthmatic airway obstruction typically results from bronchospasms. Asthma may be

triggered by a variety of causes such as allergic reactions, a secondary response to infections, industrial or occupational exposures, ingestion of certain chemicals or drugs, exercise, and vasculitis. Much of asthma's pathology can be attributed to mast cell degranulation. Mast cells will degranulate in response to various 5 conditions such as, for example, classical IgE-antigen stimulation. It is believed that when the asthmatic, human or animal, inhales an allergenic substance, sensitized IgE antibodies trigger mast cell degranulation in the lung interstitium. The mast cell degranulation releases histamine, bradykinin, and slow-reacting substance of anaphylaxis (SRS-A) which includes the leukotrienes C, D and E, 10 prostaglandins including PGF₂, PGF_{2a}, and PGD₂, and thromboxane A₂. The histamine then attaches to receptor sites in the larger bronchi, causing irritation, inflammation, and edema. The SRS-A attaches to receptor sites in the smaller bronchi, causing edema and attracting prostaglandins, which enhance the effects 15 of histamine in the lungs. Histamine, in combination with prostaglandins, also stimulates excessive mucous secretion, narrowing the bronchial lumen further. When an asthmatic individual inhales, the narrowed bronchial lumen still expands slightly, allowing air to reach the alveoli. However, upon exertion to exhale, the increased thoracic pressure closes the bronchial lumen completely. Therefore, air can enter the lungs, but may not exit during an asthma attack. The ventilation 20 in the alveoli is then inhibited by mucous collecting in the lung bases. In an effort to compensate for lowered alveolar ventilation, blood is shunted to other alveoli. Hypoxia, and in extreme cases, respiratory acidosis may result without medical intervention. In many cases, there are two phases to an allergic asthma attack,

an early phase and a late phase which follows 4-6 hours after bronchial stimulation. The early phase includes the immediate inflammatory response including the reactions caused by the release of cellular mediators from mast cells (i.e., histamine). Late phase reactions develop over a period of hours and

5 are characterized histologically by an early influx of polymorphonuclear leukocytes and fibrin deposition, later followed by infiltration of eosinophils. Increased levels of eosinophil-derived inflammatory mediators in plasma and BAL, including eosinophilic cationic protein and major basic protein, have been observed during the late phase reaction. Upregulation of TH2 -type cytokines

10 (IL4, IL5 and IL 13) following allergen challenge has also been observed during the late phase. Thus, the cellular inflammatory response, in combination with released pro-inflammatory mediators (e.g., mmp9) and locally produced cytokines in the bronchial mucosa, play a central role in the late phase allergic inflammation and bronchoconstriction. Late phase reactions increase airway

15 reactivity and lead to prolonged asthmatic exacerbations that may last from hours to days to months in some subjects. One of the residual effects of asthma reactions is this hyperresponsiveness of the airways to nonspecific stimuli.

Currently, the treatments for asthma are not always adequate and many have serious side effects. The general goals of drug therapy for asthma are

20 prevention of bronchospasm and control of airway hyperreactivity or hyperresponsiveness, an indication of airway inflammation. It is very difficult to eliminate or prevent exposure to all allergens that may trigger an asthma attack.

To prevent these attacks, most asthmatics are treated with various pharmacological agents, many of which have side effects.

In a study reported by Lukacs et al. (Lukacs et al, Am. J. Physiol Lung Cell Mol Physiol. 2001), anti-C5a antibodies were administered intratracheally

5 together with anti-BSA antibody during the induction of immune complex mediated lung inflammation. Specifically, the Lukacs study uses a model of acute immune complex mediated tissue lung inflammation similar to reverse passive Arthus reaction in skin, with a brief AHR to intravenous challenges of methacholine. Animals did not develop chronic airway inflammation featured by

10 eosinophilia; nor did they experience previous severe asthmatic attack after exposed to allergens as demonstrated in the current invention. The key feature of Lukacs study was the formation of immune complexes of BSA and anti-BSA locally along the airway, which activate the complement cascade and produced significantly amount of terminal complement components after intratracheally

15 injected anti-BSA antibody into animals. The subsequent development of AHR to methacholine lasting up to four hours during the acute phase BSA-anti-BSA induced lung inflammation, which is in significantly contrast to the severe and long lasting AHR seen patients with asthma. It is reasonable to assume based on the Lukacs study that anti-C5a antibody neutralizes C5a produced locally in

20 the airway and therefore prevents the development of harmful downstream events mediated by C5a such as recruiting and activating of inflammatory cells, stimulating the release of mediators and vascular leaky syndromes. The combination effects of blocking C5a mediated recruitment of inflammatory cells,

release of mediators such as histamine, and the development of edema of airways that leads to the reduction of airway resistance and prevention of the development of AHR when animals were challenged with IV methacholine. This study may suggest the importance of C5a in development of AHR of immune complexes mediated lung inflammation. However, this study provides no basis to predict by if there is any direct and immediate effect of bronchial dilation by C5 inhibitors during an on-going asthmatic attack or late phase airway responses to allergens. Nor does this study involve the treatment of subjects that have established airway inflammation or subjects that have experienced previous 5 asthmatic symptoms.

10

SUMMARY

The present disclosure relates to a treatment for asthma using a compound which binds to or otherwise blocks the generation and/or activity of one or more complement components or blocks the engagement of complement 15 component receptors, such as, for example, C5a receptors. The treatment therapy includes the administration of a compound that inhibits the production and/or activity of at least one complement component. Suitable compounds include, for example, antibodies which bind to or otherwise block the generation and/or activity of one or more complement components, such as, for example, an 20 antibody specific to complement component C5.

The complement-inhibiting compound can be administered prophylactically in known asthmatic individuals (such as those having established airway inflammation or a subject that has experienced previous asthmatic

symptoms) to prevent, or help prevent asthma attacks. This prophylactic therapy can be administered via intravenous, aerosol, subcutaneous or intramuscular routes.

The complement-inhibiting compound can be administered as a 5 therapeutic regimen to an individual experiencing an asthma attack. The regimen can be administered via intravenous, aerosol, subcutaneous or intramuscular routes.

A combination therapy may also be used that includes a complement-inhibiting compound in combination with a regimen of known asthma therapy, 10 such as, for example, steroids, anti-IgE antibodies, anti-IL-4 antibodies, anti-IL-5 antibodies, β 2 receptor agonists, leukotriene inhibitors, 5 Lipoxygenase inhibitors, β 2 adreno receptor agonists, PDE inhibitors, IL 5 antagonists, CD23 antagonists, IL 13 antagonists, cytokine release inhibitors, histamine H1 receptor antagonists, anti-histamines and histamine release inhibitors. Suitable 15 compounds of each class listed above as well as other asthma treatments are listed in *Asthma Therapeutic: New Treatment Options and Emerging Drug Discovery* Tasrgerts, Barnes, April 2003, Lead Discovery, <http://www.leaddiscovery.co.uk/target-discovery/abstracts/dossier-asthma.html>

In another aspect, a method of reducing inflammation in the lungs of 20 asthma patients is described herein. The methods include the step of administering to a subject having or susceptible to asthma a compound that reduces the release or production of inflammatory mediators (such as, for example, matrix metalloprotease 9 (mmp9 - also known as the 92-kDa type IV

collagenase/gelatinase or gelatinase B), TGF β , eosinophil granules, and the like) in the airways of the subject. The compound can act at the cellular level to reduce the production or release of the inflammatory mediator, can interact with the inflammatory mediator in a manner that interfaces with its activity, (such as, 5 for example by preventing the conversion of pro-mmp-9 to the 83 kDa active form or can interact with an active form) of the inflammatory mediator to prevent the inflammatory effects associated therewith.

BRIEF DESCRIPTION OF DRAWINGS

Figure 1 graphically shows the OVA-induced asthmatic reactions in 10 normal BALB/c Mice.

Figure 2a shows the schedule and nature of antigen challenges and prophylactic treatment of normal BALB/c Mice.

Figure 2b shows a schedule of antigen challenge and prophylactic treatment in normal BALB/c Mice.

15 Figure 3 graphically summarizes the effects of the treatments shown in Figures 2a and 2b.

Figure 4a shows the schedule and nature of antigen challenges and IV or aerosol treatment of normal BALB/c Mice during an asthma attack.

20 Figure 4b depicts the protocol and results of induction of asthmatic attack in BALB/c Mice.

Figures 5a and 5b graphically summarize the effects of the aerosol treatments shown in Figure 4a.

Figures 6a and 6b graphically summarize the effects of the intravenous treatments shown in Figure 4a.

Figure 7 graphically depicts the systemic effects of various treatments on C5 activity for treatments shown in Figure 4a.

5 Figure 8 graphically shows the effect of various intravenous treatments on the total WBC count in BAL.

Figures 9 and 10 show that eosinophils were found to be the predominant inflammatory cells in BAL.

10 Figure 11 graphically shows the effect of various intravenous treatments on histamine and in BAL.

Figure 12 graphically shows the effect of various intravenous treatments on MMP-9 levels in BAL.

Figure 13 graphically shows the effect of various intravenous treatments on TGF β levels in BAL.

15 Figure 14 shows the schedule and nature of antigen challenges and cannulation and aerosol treatment of normal BALB/c Mice during an asthma attack.

Figure 15 shows lung resistance in asthmatic mice treated during an asthma attack with anti-C5, β 2 receptor agonist or a combination thereof.

20 DETAILED DESCRIPTION

The present disclosure is directed to a method of treating asthma in mammals. Specifically, the methods of treating asthma described herein involve using compounds which bind to or otherwise block the generation and/or activity

of one or more complement components. The inhibition or blocking of the generation of complement components inhibits multiple factors involved in the broncho-constrictive responses in asthma. A specific class of such compounds which are particularly useful are antibodies specific to a human complement component, especially anti-C5 antibodies.

5 The complement system acts in conjunction with other immunological systems of the body to defend against intrusion of cellular and viral pathogens. There are at least 25 complement proteins, which are found as a complex collection of plasma proteins and membrane cofactors. The plasma proteins 10 make up about 10% of the globulins in vertebrate serum. Complement components achieve their immune defensive functions by interacting in a series of intricate but precise enzymatic cleavage and membrane binding events. The resulting complement cascade leads to the production of products with opsonic, immunoregulatory, and lytic functions. A concise summary of the biologic 15 activities associated with complement activation is provided, for example, in The Merck Manual, 16th Edition.

The complement cascade progresses via the classical pathway or the alternative pathway. These pathways share many components, and while they differ in their initial steps, they converge and share the same "terminal 20 complement" components (C5 through C9) responsible for the activation and destruction of target cells.

The classical complement pathway is typically initiated by antibody recognition of and binding to an antigenic site on a target cell. The alternative

pathway is usually antibody independent, and can be initiated by certain molecules on pathogen surfaces. Additionally, the lectin pathway is typically initiated with binding of mannose-binding lectin (MBL) to high mannose substrates. These pathways converge at the point where complement 5 component C3 is cleaved by an active protease (which is different in each pathway) to yield C3a and C3b. Other pathways activating complement attack can act later in the sequence of events leading to various aspects of complement function.

C3a is an anaphylatoxin. C3b binds to bacterial and other cells, as well as 10 to certain viruses and immune complexes, and tags them for removal from the circulation. (C3b in this role is known as opsonin.) The opsonic function of C3b is generally considered to be the most important anti-infective action of the complement system. Patients with genetic lesions that block C3b function are prone to infection by a broad variety of pathogenic organisms, while patients with 15 lesions later in the complement cascade sequence, i.e., patients with lesions that block C5 functions, are found to be more prone only to *Neisseria* infection, and then only somewhat more prone.

C3b also forms a complex with other components unique to each pathway to form classical or alternative C5 convertase, which cleaves C5 into C5a and 20 C5b. C3 is thus regarded as the central protein in the complement reaction sequence since it is essential to both the alternative and classical pathways. This property of C3b is regulated by the serum protease Factor I, which acts on

C3b to produce iC3b. While still functional as opsonin, iC3b cannot form an active C5 convertase.

C5 is a 190 kDa beta globulin found in normal serum at approximately 75 µg/ml (0.4 µM). C5 is glycosylated, with about 1.5-3 percent of its mass 5 attributed to carbohydrate. Mature C5 is a heterodimer of a 999 amino acid 115 kDa alpha chain that is disulfide linked to a 656 amino acid 75 kDa beta chain. C5 is synthesized as a single chain precursor protein product of a single copy gene (Haviland et al. J. Immunol. 1991, 146:362-368). The cDNA sequence of the transcript of this gene predicts a secreted pro-C5 precursor of 1659 amino 10 acids along with an 18 amino acid leader sequence (see, U.S. patent 6,355,245).

The pro-C5 precursor is cleaved after amino acid 655 and 659, to yield the beta chain as an amino terminal fragment (amino acid residues +1 to 655 of the above sequence) and the alpha chain as a carboxyl terminal fragment (amino acid residues 660 to 1658 of the above sequence), with four amino acids (amino 15 acid residues 656-659 of the above sequence) deleted between the two.

C5a is cleaved from the alpha chain of C5 by either alternative or classical C5 convertase as an amino terminal fragment comprising the first 74 amino acids of the alpha chain (i.e., amino acid residues 660-733 of the above sequence). Approximately 20 percent of the 11 kDa mass of C5a is attributed to 20 carbohydrate. The cleavage site for convertase action is at, or immediately adjacent to, amino acid residue 733 of the above sequence. A compound that would bind at, or adjacent, to this cleavage site would have the potential to block

access of the C5 convertase enzymes to the cleavage site and thereby act as a complement inhibitor.

C5 can also be activated by means other than C5 convertase activity. Limited trypsin digestion (Minta and Man, J. Immunol. 1977, 119:1597-1602;

5 Wetsel and Kolb, J. Immunol. 1982, 128:2209-2216) and acid treatment (Yammamoto and Gewurz, J. Immunol. 1978, 120:2008; Damerau et al., Molec. Immunol. 1989, 26:1133-1142) can also cleave C5 and produce active C5b.

Cleavage of C5 releases C5a, a potent anaphylatoxin and chemotactic factor, and leads to the formation of the lytic terminal complement complex, C5b-

10 9. C5a and C5b-9 also have pleiotropic cell activating properties, by amplifying the release of downstream inflammatory factors, such as hydrolytic enzymes, reactive oxygen species arachidonic acid metabolites and various cytokines.

C5b combines with C6, C7, and C8 to form the C5b-8 complex at the surface of the target cell. Upon binding of several C9 molecules, the membrane

15 attack complex (MAC, C5b-9, terminal complement complex--TCC) is formed.

When sufficient numbers of MACs insert into target cell membranes the openings they create (MAC pores) mediate rapid osmotic lysis of the target cells. Lower, non-lytic concentrations of MACs can produce other effects. In particular,

membrane insertion of small numbers of the C5b-9 complexes into endothelial 20 cells and platelets can cause deleterious cell activation. In some cases activation may precede cell lysis.

As mentioned above, C3a and C5a are anaphylatoxins. These activated complement components can trigger mast cell degranulation, which releases

histamine from basophils and mast cells, and other mediators of inflammation, resulting in smooth muscle contraction, increased vascular permeability, leukocyte activation, and other inflammatory phenomena including cellular proliferation resulting in hypercellularity. C5a also functions as a chemotactic 5 peptide that serves to attract pro-inflammatory granulocytes to the site of complement activation.

C5a receptors are found on the surfaces of bronchial and alveolar epithelial cells and bronchial smooth muscles cells. C5a receptors have also been found on eosinophils, mast cells, monocytes, neutrophils, and activated 10 lymphocytes. Thus, compounds that block engagement of receptors of complement components are useful herein.

Any compounds which bind to or otherwise block the generation and/or activity of any of the human complement components, such as, for example, antibodies specific to a human complement component are useful herein. Some 15 compounds include antibodies directed against complement components C-1, C-2, C-3, C-4, C-5, C-6, C-7, C-8, C-9, Factor D, Factor B, Factor P, MBL, MASP-1, AND MASP-2, thus preventing the generation of the anaphylatoxic activity associated with C5a and preventing the assembly of the membrane attack complex associated with C5b. Also useful in the present methods are naturally 20 occurring or soluble forms of complement inhibitory compounds such as CR1, LEX-CR1, MCP, DAF, CD59, Factor H, cobra venom factor, FUT-175, y bind protein, complestatin, and K76 COOH.

Particularly useful compounds for use herein are antibodies that reduce, directly or indirectly, the conversion of complement component C5 into complement components C5a and C5b. One class of useful antibodies are those having at least one antibody-antigen binding site and exhibiting specific

5 binding to human complement component C5, wherein the specific binding is targeted to the alpha chain of human complement component C5. More particularly, a monoclonal antibody (mAb) may be used. Such an antibody 1) inhibits complement activation in a human body fluid; 2) inhibits the binding of purified human complement component C5 to either human complement

10 component C3 or human complement component C4; and 3) does not specifically bind to the human complement activation product for C5a.

Particularly useful complement inhibitors are compounds which reduce the generation of C5a and/or C5b-9 by greater than about 30%. Anti-C5 antibodies that have the desirable ability to block the generation of C5a have been known in

15 the art since at least 1982 (Moongkarndi et al. Immunobiol. 1982, 162:397; Moongkarndi et al. Immunobiol. 1983, 165:323). Antibodies known in the art that are immunoreactive against C5 or C5 fragments include antibodies against the C5 beta chain (Moongkarndi et al. Immunobiol. 1982, 162:397; Moongkarndi et al. Immunobiol. 1983, 165:323; Wurzner et al. 1991, *supra*; Mollnes et al. Scand. J. Immunol. 1988, 28:307-312); C5a (see for example, Ames et al. J. Immunol. 1994, 152:4572-4581, U.S. Pat. No. 4,686,100, and European patent publication No. 0 411 306); and antibodies against non-human C5 (see for example, Giclas et al. J. Immunol. Meth. 1987, 105:201-209). Particularly useful anti-C5

antibodies are h5G1.1, h5G1.1-scFv and functional fragments of h5G1.1.

Methods for the preparation of h5G1.1, h5G1.1-scFv and functional fragments of h5G1.1 are described in U.S. Patent No. 6,355,245 and "Inhibition of Complement Activity by Humanized Anti-C5 Antibody and Single Chain Fv",

5 Thomas et al., *Molecular Immunology*, Vol. 33, No. 17/18, pages 1389-1401, 1996, the disclosures of which are incorporated herein in their entirety by this reference.

Functionally, a suitable antibody inhibits the cleavage of C5, which blocks the generation of potent proinflammatory molecules C5a and C5b-9 (terminal 10 complement complex). Preferably, the antibody does not prevent the formation of C3b, which subserves critical immunoprotective functions of opsonization and immune complex clearance.

While preventing the generation of these proinflammatory terminal complement components, antibody-mediated inhibition of the complement 15 cascade at C5 preserves the ability to generate C3b, which is critical for opsonization of many pathogenic microorganisms, as well as for immune complex solubilization and clearance. Retaining the capacity to generate C3b appears to be particularly important as a therapeutic factor in complement inhibition for inflammatory diseases, where increased susceptibility to infection 20 and impaired clearance of immune complexes are preexisting clinical features of the disease process.

The anti-human C5 antibody is preferably a monoclonal antibody, although polyclonal antibodies produced and screened by conventional techniques can also be used if desired.

The preferred anti-C5 antibodies used to treat asthma in accordance with 5 this disclosure bind to C5 or fragments thereof, e.g., C5a or C5b. Preferably, the anti-C5 antibodies are immunoreactive against epitopes on the alpha chain of purified human complement component C5 and are capable of blocking the conversion of C5 into C5a and C5b by C5 convertase. This capability can be measured using the techniques described in Wurzner, et al., Complement 10 Inflamm 8:328-340, 1991.

In a particularly useful embodiment, the anti-C5 antibodies are not immunoreactive against epitopes on the beta chain, but rather are immunoreactive against epitopes within the alpha chain of purified human complement component C5. In this embodiment, the antibodies are also capable 15 of blocking the conversion of C5 into C5a and C5b by C5 convertase. Within the alpha chain, the most preferred antibodies bind to an amino-terminal region, however, they do not bind to free C5a.

Hybridomas producing monoclonal antibodies reactive with complement component C5 can be obtained according to the teachings of Sims, et al., U.S. 20 Pat. No. 5,135,916. Antibodies are prepared using purified components of the complement membrane attack complex as immunogens according to known methods. In accordance with this disclosure, complement component C5 or C5b

is preferably used as the immunogen. In accordance with particularly preferred useful embodiments, the immunogen is the alpha chain of C5.

Particularly useful antibodies share the required functional properties discussed in the preceding paragraph and have any of the following

5 characteristics:

(1) they compete for binding to portions of C5—the C5 alpha chain; and

(2) they specifically bind to the C5 alpha chain. Such specific binding, and competition for binding can be determined by various methods well known in the art, including the plasmon surface resonance method (Johne et al., *J. Immunol.*

10 *Meth. 1993, 160:191-198*).

(3) they block the binding of C5 to either C3 or C4 (which are components of C5 convertase).

In another aspect, a method of reducing inflammation in the lungs of asthma patients is provided. The methods include the step of administering to a 15 subject having or susceptible to asthma a compound that reduces the production or release of inflammatory mediators in the airway of the subject. Non-limiting examples of inflammatory mediators that can be reduced in accordance with this disclosure include TGF β , eosinophil granule proteins, and matrix metalloprotease 9 (mmp9 - also known as the 92-kDa type IV collagenase/gelatinase or 20 gelatinase B). The compounds can reduce inflammation by any variety of mechanisms. Where the inflammatory mediator is MMP-9, for example, the compound can act at the cellular level to reduce the production or release of pro-mmp-9, can interact with pro-mmp-9 in a manner that prevents conversion of pro-

mmp-9 to the 83 kDa active form or can interact with the active form of mmp9 to prevent the inflammatory effects associated with the enzyme (such as, for example, the generation of TGF-beta). Suitable compounds include, but are not limited to, the compounds described above which bind to or otherwise block the 5 generation and/or release an/or activity of one or more complement components or block engagement of the receptors of complement components.

MMP-9 activity can be detected in accordance with a variety of art-recognized procedures. For example, quantitative zymographic methods provide a relatively refined assessment of the activity of this enzyme. This method allows 10 for the detection of MMP-9 activity based upon the ability of the enzyme to hydrolyze denatured collagen, i.e., gelatin, which is a natural substrate for MMP-9. The gelatin is incorporated into a gel such as polyacrylamide. See Hibbs et al., J. Biol. Chem. 260:2493-2500 (1985) and Moll et al., Cancer Res. 50:6162-70 (1990). The assay may be standardized using a purified MMP-9 preparation that 15 is analyzed in parallel with the test sample. Purified MMP-9 can be prepared by methods known in the art. See, for example, Okada et al., *supra*., and Morodomi et al., Biochem J. 285:603:11 (1992). The extent of hydrolysis of the gelatin is directly related to the activity of MMP-9 in the sample, and the active MMP-9 forms can be identified by their characteristic molecular weights. In the gelatin 20 zymography, the proMMP-9 species can be detected because of the catalytic activation that occurs during electrophoresis and subsequent incubation. However, the MMP-9 forms present prior to the onset of labor are incapable of undergoing this kind of activation, i.e., they are latent.

MMP-9 activity can also be detected using standard immunological techniques, e.g., ELISA, immunofluorescence assays, or radioimmunoassays. In a preferred embodiment, MMP-9 activity is detected using ELISA, which entails the use of antibodies specific to MMP-9. See David et al, U.S. Pat. No. 4,376,110 (and references cited therein). Monoclonal antibodies specific to MMP-9 have been prepared using partially purified enzyme preparations. See, e.g., Moll et al., *supra*; Ramos-DeSimone et al., *HYBRIDOMA* 12(4):349-63 (1993) and Goldberg et al. Polyclonal antibodies specific to MMP-9 can also be prepared in accordance with standard procedures. In a preferred embodiment, polyclonal antibodies are prepared using non-conserved peptides conjugated to a macromolecular carrier. The choice of a specific non-conserved peptide such as the metal-binding domain, among the members of the MMPs is considered within a level of ordinary skill in the art. See Woessner, and Goldberg et al., *supra*. Enzymic assays that can detect MMP-9 in picogram or nanogram amounts are also disclosed in Manicourt et al., *Anal. Biochem.* 215(2):171-9 (1993).

MMP-9 activity can further be detected in a sample by western blot analysis, which requires electrophoretic separation of the test material in a gel, followed by transfer of the separated proteins to a nitrocellulose membrane and detection of the MMP-9 antigens with a specific antibody and reagent that reacts with the antigen-fixed antibody. See Towbin et al., *Proc. Natl. Acad. Sci. USA* 76(9):4350-4354 (1979).

Suitable assays for detection of mmp9 are commercially available from a variety of sources such as, for example Boehringer Manheim Biochemicals (Manheim, Germany) and R&D Systems, (Minneapolis, MN).

The compound that inhibits the production and/or activity of at least one 5 complement component can be administered in a variety of unit dosage forms.

The dose will vary according to the particular compound employed. For example, different antibodies may have different masses and/or affinities, and thus require different dosage levels. Antibodies prepared as Fab' fragments will also require differing dosages than the equivalent intact immunoglobulins, as 10 they are of considerably smaller mass than intact immunoglobulins, and thus require lower dosages to reach the same molar levels in the patient's blood.

The dose will also vary depending on the manner of administration, the particular symptoms of the patient being treated, the overall health, condition, size, and age of the patient, and the judgment of the prescribing physician.

15 Administration of the compound that inhibits the production and/or activity of at least one complement component will generally be in an aerosol form with a suitable pharmaceutical carrier, via intravenous infusion by injection, or subcutaneous injection. Other routes of administration may be used if desired. Aerosol administration is preferred since it avoids the systemic effects of the 20 complement-inhibiting compound, while providing the desired asthma-treating effects to be achieved in accordance with this disclosure.

Formulations suitable for injection are found in Remington's Pharmaceutical Sciences, Mack Publishing Company, Philadelphia, Pa., 17th ed.

(1985). Such formulations must be sterile and non-pyrogenic, and generally will include a pharmaceutically effective carrier, such as saline, buffered (e.g., phosphate buffered) saline, Hank's solution, Ringer's solution, dextrose/saline, glucose solutions, and the like. The formulations may contain pharmaceutically acceptable auxiliary substances as required, such as, tonicity adjusting agents, wetting agents, bactericidal agents, preservatives, stabilizers, and the like.

EXAMPLE 1-The Use Of Anti-C5 Antibodies As A Prophylactic For C5 Inhibition To Treat Asthma.

10 Asthma was induced in normal BALB/c mice by exposing them to Ovalbumin antigen ("OVA") and Alum, according to the dosages and schedule shown in Figure 4b. These exposures solicited the standard asthmatic response during an attack. The exposed mice showed both early phase and late phase reactions. As seen in Figure 1, the early phase reaction was specific airway 15 resistance within 15 minutes after exposure. A more severe late phase reaction of specific airway resistance occurred approximately 5 hours after exposure. Specific airway resistance was measured by means of double chamber plethysmograph fitted with pneumotachographs, which is commercially available from the Buxo Corporation.

20 To demonstrate the prophylactic benefits of a complement component inhibiting compound, groups of mice were given one of three different treatments in accordance with the schedule of dosages shown in Figures 2a and 2b. The positive control group was treated with a control antibody (hybridoma 135.8) at a dose of 40 mg/kg. A second group of mice was treated with the anti-C5 25 antibody, BB5.1, at a dose of 40mg/kg. A third group of mice was treated with

Dexamethasone ("DEX"). A negative control group was initially exposed to PBS (phosphate buffer saline) and Alum, and was later given PBS as shown in the schedule of Figure 2a. This control group provided a baseline air resistance measurement.

5 The BB5.1 antibody is made according to known methods. (See, Frei, Y., Lambris, J.D., Stockinger, B. Mol. Cell. Probes. 1: 141-149 (1987)). Both the BB5.1 antibody and the isotype match control 135.8 hybridoma antibody were grown as ascites in athymic mice and the antibodies were purified from ascites by protein A affinity chromatography followed by elution with ImmunoPure IgG 10 elution buffer (Pierce) and dialysis against PBS buffered saline. (Wang et al. 1996).

It was found that the group prophylactically treated subcutaneously with Anti-C5 antibody responded as well as the challenged mice treated with steroids and the negative control, as shown in Figure 3. The Anti-C5 treatments had 15 inhibited the production of complement component C5 in the cascade, and by inhibiting C5 production, the mice experienced significantly less airway constriction. The positive control group that was treated with a control antibody showed increased specific airway resistance caused by an asthmatic attack.

EXAMPLE 2 - The Use Of Anti-C5 Antibodies Via Intravenous Administration As A Therapeutic Method To Treat Asthma

To demonstrate the therapeutic benefits of administering a C5

5 complement component inhibitor intravenously during an asthma attack, groups of mice were given one of three different treatments in accordance with the schedule of dosages shown in Figure 4a. A positive control group was challenged with OVA and treated with a control antibody (135.8 hybridoma) administered intravenously, 15 minutes after provoking the initial asthmatic 10 attack. A second group of mice were similarly challenged and intravenously treated with BB5.1, the anti-C5 antibody, 15 minutes after provoking the initial asthmatic attack. A third group of mice were similarly challenged and intravenously treated with DEX, 15 minutes after provoking the initial asthmatic attack. Sham operated mice provided baseline air resistance measurement 15 which was provided by administering phosphate buffer solution (PBS) and alum to a group of mice, and administering aerosol doses of PBS according to the schedule shown on Fig. 4a.

The results of this challenge (shown in Figs. 6a and 6b) found that the group of mice treated therapeutically with anti-C5 intravenously showed very little 20 specific airway constriction and these mice responded to the treatments as well as the group treated with the steroid, DEX. The positive control group that was treated with a control antibody showed significantly increased specific airway resistance, as shown in Figure 6a. The anti-C5 antibody had inhibited the

inflammatory response of the complement components, and allowed greater air passage during the asthma attacks in the mice.

EXAMPLE 3 - The Use Of Anti-C5 Antibodies Via Aerosol Administration As A Therapeutic Method To Treat Asthma

5

To demonstrate the therapeutic benefits of administering a C5 complement component inhibitor via aerosol during an asthma attack, groups of mice were given one of three different treatments in accordance with the schedule of dosages shown in Figure 4a. A positive control group was 10 challenged with OVA and treated with a control antibody (135.8 hybridoma) administered by aerosol, 15 minutes after provoking the initial asthmatic attack. A second group of mice were similarly challenged and treated via aerosol administration of BB5.1, an anti-C5 antibody, 15 minutes after provoking the initial asthmatic attack. A third group of mice were similarly challenged and 15 treated by aerosol with DEX, 15 minutes after provoking the initial asthmatic attack. A baseline air resistance measurement was provided by administering phosphate buffer solution (PBS) and alum to a group of mice, and administering aerosol doses of PBS according to the schedule shown on Fig. 4a.

It was found that therapeutic aerosol treatments of mice with anti-C5 20 antibody, BB5.1 during asthma attacks significantly lowered the specific airway resistance and these mice responded as well as, and more quickly than, the steroid treated mice. The results of the aerosol treatments are shown in Figures 5a and 5b. The positive control group that was treated with a control antibody

experienced specific airway resistance many times greater than the mice treated with anti-C5 antibody, as shown in Figure 5b.

Anti-C5 antibody, BB5.1 was also administered to the test animals through nebulization, which was found to be an effective method of administration.

5 Results of serum tests after nebulization indicate that the BB5.1 was binding with the C5 site and, therefore, remained intact during delivery through nebulization.

The systemic effect of each treatment given in Examples 2 and 3 was measured using the techniques described in Wurzner, et al., Complement Inflamm 8:328-340, 1991 for hemolytic activity. The results of these tests are 10 shown in Figure 7. As seen therein, the control antibody and steroid did not substantially reduce systemic C5 activity, irrespective of the method of administration. With the anti-C5 antibody BB5.1, however, the manner of administration directly affected systemic C5 activity. Specifically, although both aerosol and intravenous administration were effective at reducing the severity of 15 an asthma attack, the aerosol administration did so without substantially reducing systemic C5 activity. As seen in Figure 7, intravenous administration of the anti-C5 antibody reduced systemic C5 activity by nearly 80%.

EXAMPLE 4 - The Use Of Anti-C5 Antibodies To Reduce the
Presence of Inflammatory mediators

20

To demonstrate the therapeutic benefits of administering a C5 complement component inhibitor in reducing the presence of inflammatory mediators during an asthma attack, groups of mice were given one of three different treatments in accordance with the schedule of dosages shown in Figure

4a. A positive control group was challenged with OVA and treated with a control antibody (135.8 hybridoma) administered intravenously, 15 minutes after provoking the initial asthmatic attack. A second group of mice were similarly challenged and intravenously treated with BB5.1, the anti-C5 antibody, 15

5 minutes after provoking the initial asthmatic attack. A third group of mice were similarly challenged and intravenously treated with DEX, 15 minutes after provoking the initial asthmatic attack. Sham operated mice provided baseline air resistance measurement which was provided by administering phosphate buffer solution (PBS) and alum to a group of mice, and administering aerosol

10 doses of PBS according to the schedule shown on Fig. 4a. After 5 hours the mice were euthanized and the lungs were lavaged using conventional techniques. Generally, 1 cc of PBI saline was introduced into the lungs and recovered. This process was repeated three times. The fluid was recovered and centrifuged. The cells contained in the resulting pellet were inspected. The

15 supernatant was tested for the presence of histamine, IL-5, IL-4, IL-13, Eosinophils granule proteins, TGF β and/or mmp-9. The assays were all conducted using commercially available test kits. The results are shown in Figures 8 through 13. These BronchoAlveolarLavage (BAL) results show the presence or absence of proteins which were produced or released by lung

20 structure cells or inflammatory cells, rather than the presence of such components in the lung tissue itself.

It was found that treatment of mice with anti-C5 antibody and DEX both reduced the total WBC count. (See Fig. 8). Eosinophils were found to be the predominant inflammatory cells in the BAL. (See Figs. 9A-D and 10.)

Figure 11 shows that the anti-C5 antibody had little effect on histamine level, a result similar to that obtained with the steroid DEX. However, the anti-C5 antibody significantly reduced the level of mmp-9 detected in BAL as shown in Figure 12, compared to treatment with Steroid. Thus, while both anti-C5 antibody and DEX resulted in a lower detectable amount of TGF- β (see Fig. 13), only the anti-C5 antibody did so through a mechanism that involved a reduction in production or release of mmp-9.

EXAMPLE 5 - The Use Of Anti-C5 Antibodies As Bronchial Dilator

To demonstrate the direct and immediate bronchial dilation effect of administering a C5 complement component inhibitor (alone or in combination with a β 2 adreno receptor agonist), during an asthma attack, groups of mice were challenged with antigen and given one of four different treatments in accordance with the schedule of dosages shown in Figures 14 and 15. A positive control group was challenged with OVA, cannulated and treated with a control antibody (mouse IgG1) administered via aerosol after provoking a second asthmatic attack. A second group of mice were similarly challenged and treated with BB5.1, an anti-C5 antibody after provoking a second asthmatic attack. A third group of mice were similarly challenged and treated with salbutamol (a β 2 adreno receptor agonist commercially available from Sigma) after provoking a second asthmatic attack. A fourth group of mice were similarly challenged and treated

with a combination of anti-C5 antibody and salbutamol after provoking a second asthmatic attack. Sham operated mice provided baseline air resistance measurement which was provided by administering phosphorate buffer solution (PBS) and alum to a group of mice, and administering aerosol doses of PBS

5 according to the schedule shown on Figs. 14 and 15.

Airway responsiveness was then assessed as a change in airway function by an invasive method wherein changes in lung resistance were measured by using Buxco Biosystem software and whole body plethysmograph. Mice were anesthetized with Avertin (160mg/kg) by i.p. injection and ventilated

10 by a Harvard Apparatus Inspira ventilator. After tracheal cannula, pancuronium (0.3mg/kg) was injected intraperitoneally to induce paralysis and inhibit spontaneous breathing. The mice are kept breathing by a ventilator, which is set at a tidal volume and respiratory rate by program of body weight. The measurements of RL to specific antigen were performed at 5 hours after 5% OVA

15 provocation. The results, which are reported in Figure 15 show that the treatment with anti-C5 antibody had a significant bronchial dilation effect in 3 of 4 mice and that a synergistic effect is observed from the treatment with anti-C5 antibody in combination with a β 2 adreno receptor agonist.

Although preferred and other embodiments of the invention have been

20 described herein, further embodiments may be perceived by those skilled in the art without departing from the scope of the invention.

CLAIMS

What is claimed is:

1. A method of treating asthma in a subject comprising administering an anti-C5 antibody to a subject susceptible to or having asthma.
- 5 2. A method of preventing asthma attacks comprising prophylactically administering an anti-C5 antibody to a subject having established airway inflammation or a subject that has experienced previous asthmatic symptoms.
- 10 3. A method of reducing the severity of an asthma attack comprising administering an anti-C5 antibody to a subject having an asthma attack.
4. A method of reducing airway obstruction in a subject comprising administering an anti-C5 antibody to the subject.
- 15 5. A method of increasing air flow in a subject comprising administering an anti-C5 antibody to the subject.
6. A method of reducing bronchial spasms in a subject comprising administering an anti-C5 antibody to the subject.
- 17 7. A method of treating a chronic obstructive pulmonary disease in a subject comprising administering an anti-C5 antibody to the subject afflicted with a chronic obstructive pulmonary disease.
- 20 8. A method of reducing inflammation in a subject comprising administering an anti-C5 antibody to a subject having established airway inflammation or a subject that has experienced previous asthmatic symptoms.

9. A method of treating a subject having established airway inflammation or a subject that has experienced previous asthmatic symptoms comprising administering an effective bronchial-dilating amount of an anti-C5 antibody.

5 10. A method as in claim 8 or 9 wherein the step of administering comprises administering the anti-C5 antibody during an asthma attack.

11. A method as in any of claims 1-9 wherein the subject is a human.

12. A method as in any of claims 1-9 wherein the step of administering an anti-C5 antibody comprises administering an anti-C5 antibody that inhibits 10 the conversion of complement component C5 into C5a and C5b.

13. A method as in any of claims 1-9 wherein the step of administering an anti-C5 antibody comprises administering an anti-C5 antibody that binds to human complement component C5a.

14. A method as in any of claims 1-9 wherein the step of administering an anti-C5 antibody to human complement component C5b9.

15. A method as in any of claims 1-9 wherein the step of administering an anti-C5 antibody comprises administering an anti-C5 antibody selected from the group consisting of h5G1.1, h5G1.1-scFv and functional fragments of h5G1.1.

20 16. A method as in any of claims 1-9 wherein the step of administering an anti-C5 antibody comprises administering an anti-C5 antibody that is an antibody comprising at least one antibody-antigen binding site, said antibody exhibiting specific binding to the alpha chain of human

complement component C5, wherein the antibody 1) inhibits complement activation in a human body fluid; 2) inhibits the binding of purified human complement component C5 to C5 convertase.

17. A method as in any of claims 1-9 wherein the step of
5 administering an anti-C5 antibody comprises administering an anti-C5 antibody as an aerosol.

18. A method as in any of claims 1-9 wherein the step of administering an anti-C5 antibody comprises administering an anti-C5 antibody that via a method selected from the group consisting of intravenous infusion by
10 injection and subcutaneous injection.

19. A method as in any of claims 1-9 wherein the step of administering an anti-C5 antibody comprises administering an anti-C5 antibody in combination with a member selected from the group consisting of steroids, anti-IgE antibodies, anti-IL-4 antibodies, anti-IL-5 antibodies, β 2 adreno receptor agonists, leukotriene inhibitors, 5 Lipoxygenase inhibitors, β 2 adreno receptor agonists, PDE inhibitors, CD23 antagonists, IL 13 antagonists, cytokine release inhibitors, histamine H1 receptor antagonists, anti-histamines and histamine release inhibitors.
15

20. A method for treating a subject having or susceptible to asthma comprising administering at least one member selected from the group consisting of steroids, anti-IgE antibodies, anti-IL-4 antibodies, anti-IL-5 antibodies, β 2 adreno receptor agonists, leukotriene inhibitors, 5 Lipoxygenase inhibitors, β 2 adreno receptor agonists, PDE inhibitors,
20

CD23 antagonists, IL 13 antagonists, cytokine release inhibitors, histamine H1 receptor antagonists, anti-histamines and histamine release inhibitors in combination with an anti-C5 antibody.

21. A method of treating asthma comprising administering an anti-C5 antibody

5 to the lungs of a subject without substantially reducing systemic complement activity in the subject.

22. A method of treating asthma in a subject comprising administering a

compound to the subject, the compound being selected from the group consisting of compounds which bind to one or more complement components, compounds which block the generation of one or more complement components, compounds which block the activity of one or more complement components and compounds which block the engagement of complement component receptors.

10 23. A method as in claim 22 wherein the step of administering comprises

15 administering an anti-C5a receptor antibody.

24. A method of reducing inflammation in a subject comprising administering a

compound to the subject, the compound being selected from the group consisting of compounds which bind to one or more complement components, compounds which block the generation of one or more complement components, compounds which block the activity of one or more complement components and compounds which block the engagement of complement component receptors.

25. A method as in claim 24 wherein the step of administering comprises
administering the anti-C5 antibody during an asthma attack.

26. A method as in claim 24 wherein the step of administering comprises
administering an anti-C5a receptor antibody.

5 27. A method of treating a subject having established airway inflammation or a
subject that has experienced previous asthmatic symptoms comprising
administering a compound to the subject, the compound being selected
from the group consisting of compounds which bind to one or more
complement components, compounds which block the generation of one
10 or more complement components, compounds which block the activity of
one or more complement components and compounds which block the
engagement of complement component receptors.

28. A method as in claim 27 wherein the step of administering comprises
administering the anti-C5 antibody during an asthma attack.

15 29. A method as in claim 27 wherein the step of administering comprises
administering an anti-C5a receptor antibody.

30. A method as in any of claims 22, 24 or 27 wherein the step of
administering a compound comprises administering one or more
antibodies directed against a compound selected from the group
20 consisting of complement components C-1, C-2, C-3, C-4, C-5, C-6, C-7,
C-8, C-9, Factor D, Factor B, Factor P, MBL, MASP-1, and MASP-2.

31. A method as in any of claims 22, 24 or 27 wherein the step of
administering a compound comprises administering one or more

compounds selected from the group consisting of soluble CR1, soluble LEX-CR1, soluble MCP, soluble DAF, soluble CD59, Factor H, cobra venom factor, FUT-175, complestatin, and K76 COOH.

32. A method of preventing asthma attacks comprising prophylactically
5 administering a compound to a subject having established airway
inflammation or a subject that has experienced previous asthmatic
symptoms, the compound being selected from the group consisting of
compounds which bind to one or more complement components,
compounds which block the generation of one or more complement
components, compounds which block the activity of one or more
10 complement components and compounds which block the engagement of
complement component receptors.

33. A method as in claim 32 wherein the step of administering comprises
administering an anti-C5a receptor antibody.

15 34. A method of reducing the severity of an asthma attack comprising
administering a compound to a subject having an asthma attack, the
compound being selected from the group consisting of compounds which
bind to one or more complement components, compounds which block the
generation of one or more complement components, compounds which
block the activity of one or more complement components and compounds
20 which block the engagement of complement component receptors.

35. A method as in claim 34 wherein the step of administering comprises
administering an anti-C5a receptor antibody.

36. A method of reducing inflammation in the lungs of an asthma patient comprising the step of administering to a subject having or susceptible to asthma one or more compounds selected from the group consisting of soluble CR1, soluble LEX-CR1, soluble MCP, soluble DAF, soluble CD59, Factor H, cobra venom factor, FUT-175, complestatin, K76 COOH and antibodies directed against a compound selected from the group consisting of complement components C-1, C-2, C-3, C-4, C-5, C-6, C-7, C-8, C-9, Factor D, Factor B, Factor P, MBL, MASP-1, and MASP-2.

5 37. A method as in claim 36 wherein the compound administered reduces the release or production of one or more inflammatory mediators in the 10 airways of the subject.

38. A method as in claim 36 wherein the step of administering comprises administering a compound that acts at the cellular level to reduce the production or release of an inflammatory mediator.

15 39. A method as in claim 36 wherein the step of administering comprises administering a compound that can interact with an active form of the inflammatory mediator to prevent the inflammatory effects associated therewith.

40. A method as in claim 36 wherein the step of administering a compound 20 comprises administering one or more antibodies directed against a compound selected from the group consisting of complement components C-1, C-2, C-3, C-4, C-5, C-6, C-7, C-8, C-9, Factor D, Factor B, Factor P, MBL, MASP-1, and MASP-2.

41. A method as in claim 36 wherein the step of administering a compound comprises administering one or more compounds selected from the group consisting of CR1, LEX-CR1, MCP, DAF, CD59, Factor H, cobra venom factor, FUT-175, complestatin, and K76 COOH.

5 42. A method as in claim 36 wherein the step of administering a compound comprises administering the compound in combination with a member selected from the group consisting of steroids, anti-IgE antibodies, anti-IL-4 antibodies, anti-IL-5 antibodies, β 2 receptor agonists, leukotriene inhibitors, 5 Lipoxygenase inhibitors, β 2 adreno receptor agonists, PDE inhibitors, CD23 antagonists, IL 13 antagonists, cytokine release inhibitors, histamine H1 receptor antagonists, anti-histamines and histamine release inhibitors.

10 43. A method as in claim 36 wherein the step of administering a compound comprises administering an anti-C5 antibody.

15 44. A method as in claim 34 wherein the step of administering comprises administering an anti-C5a receptor antibody.

1/19

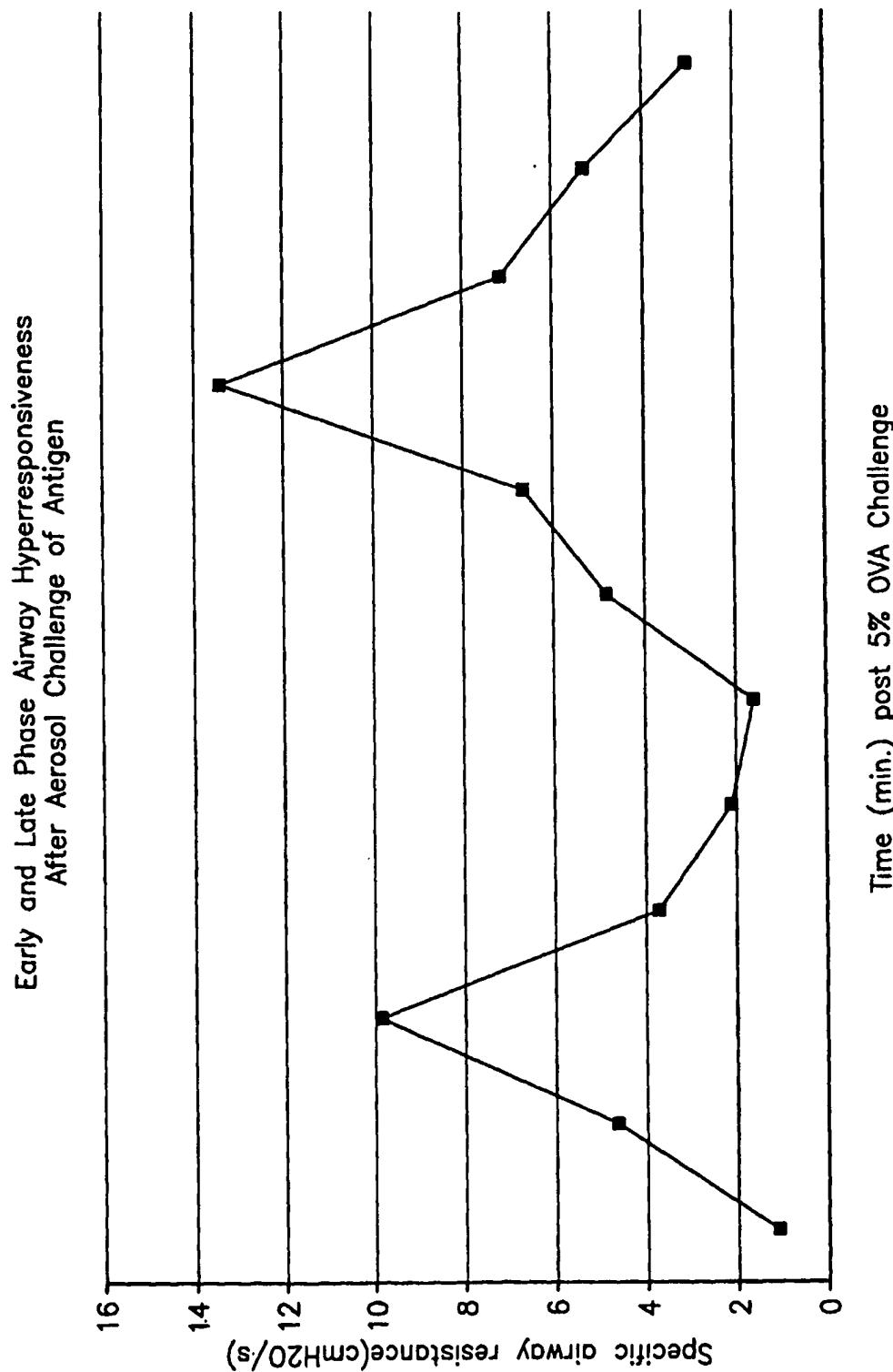


FIG. 1

2/19

Induction of Asthmatic Attack in Normal BALB/c Mice and Effect of Prophylactic C5 Inhibition

<u>Group & Schedule:</u>	<u>day 1, 14</u>	<u>day 25, 29, 32</u>	<u>day 28, 29, 30</u>	<u>day 33</u>
P. Control	Ova + Alum	135.8, 40mg/kg	1% OVA aerosol	5% OVA aerosol
Anti-C5	Ova + Alum	BB5.1, 40mg/kg	1% OVA aerosol	5% OVA aerosol
Steroid	Ova + Alum	Dex	1% OVA aerosol	5% OVA aerosol
N. Control	PBS + Alum	PBS	PBS	PBS

FIG. 2a

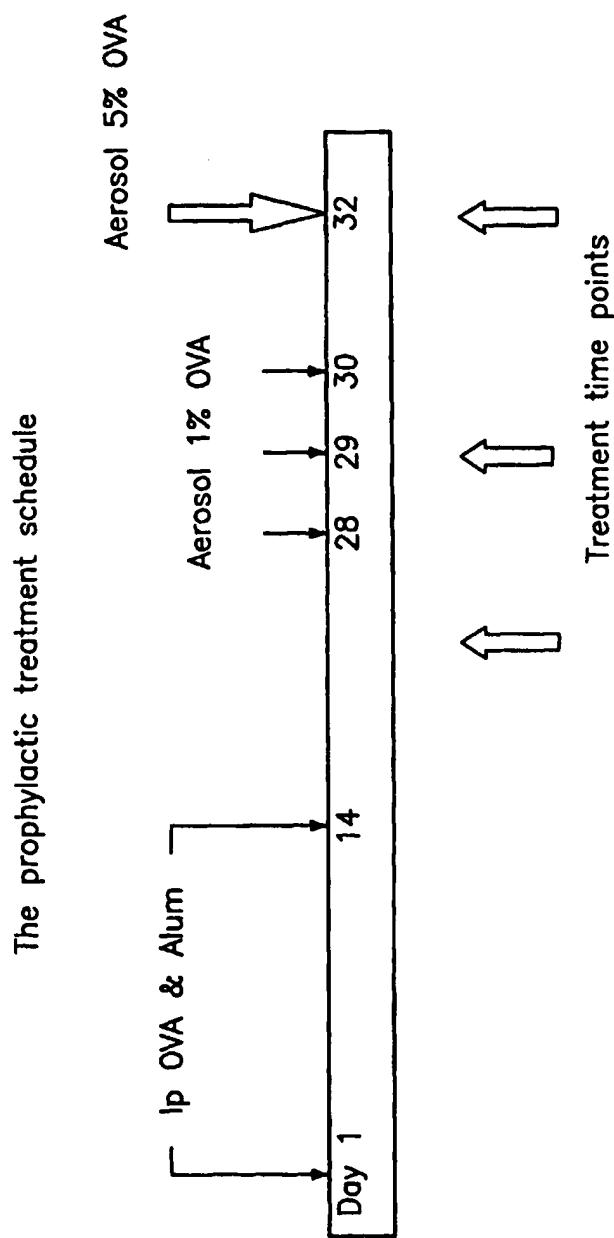


FIG. 2b

Induction of Asthmatic Attack in Normal BALB/c Mice
and Effect of Prophylactic C5 Inhibition

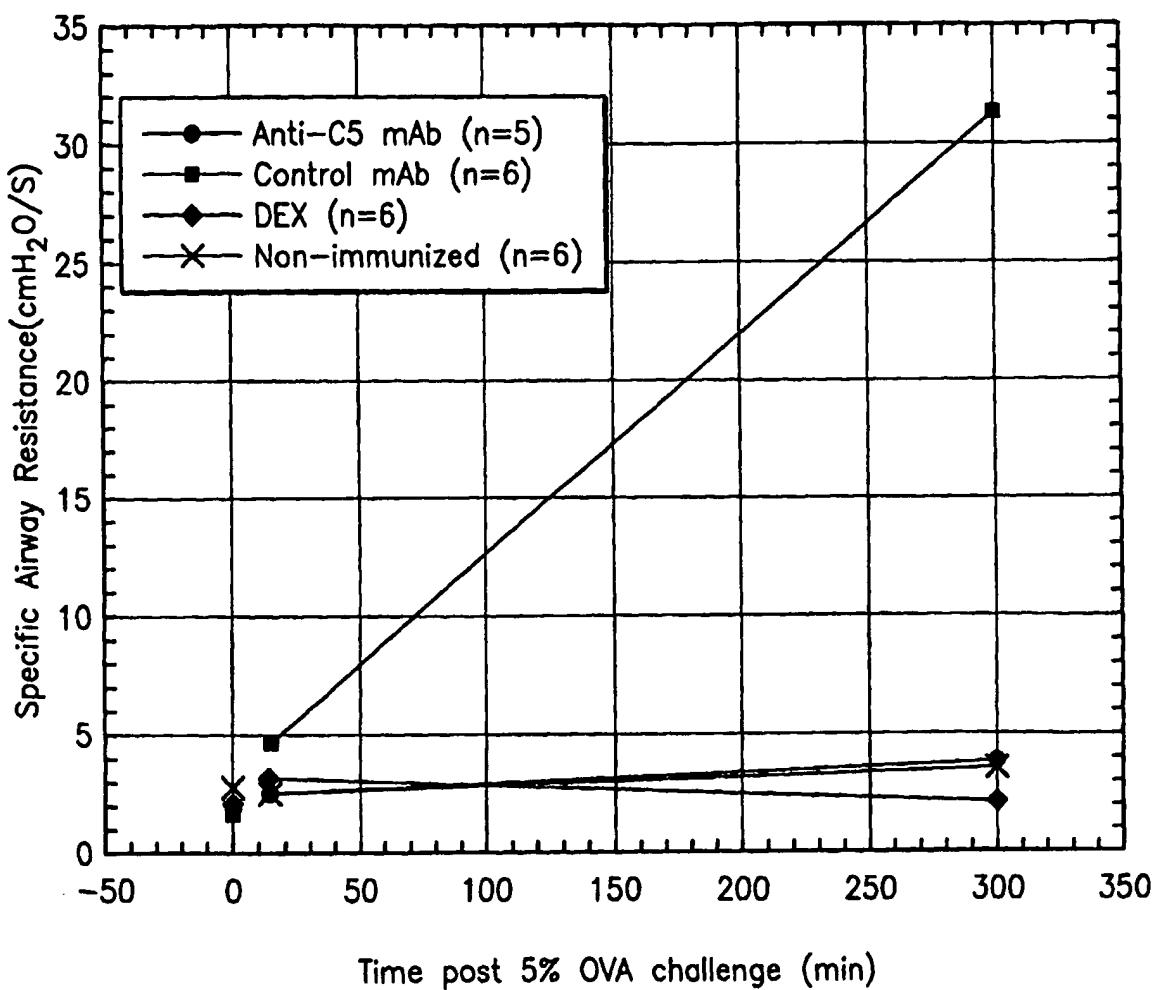


FIG. 3

5/19

Effect of Therapeutic C5 Inhibition (Aerosol or IV) During Asthmatic Attack

<u>Group & Schedule:</u>	<u>day 1, 14</u>	<u>Aerosol on day 28-30 & day 32</u>	<u>day 35</u>	<u>day 35</u>
P.Control	Ova + Alum	1% ova on day 29-30, 5% on 32	5% OVA	IV or aerosol 135.8
Anti-C5	Ova + Alum	1% ova on day 29-30, 5% on 32	5% OVA	IV or aerosol BB5.1
Steroid	Ova + Alum	1% ova on day 29-30, 5% on 32	5% OVA	IV or aerosol Dex
N.Control	PBS + Alum	PBS on day 29-30, PBS on 32	PBS	IV or aerosol PBS

FIG. 4a

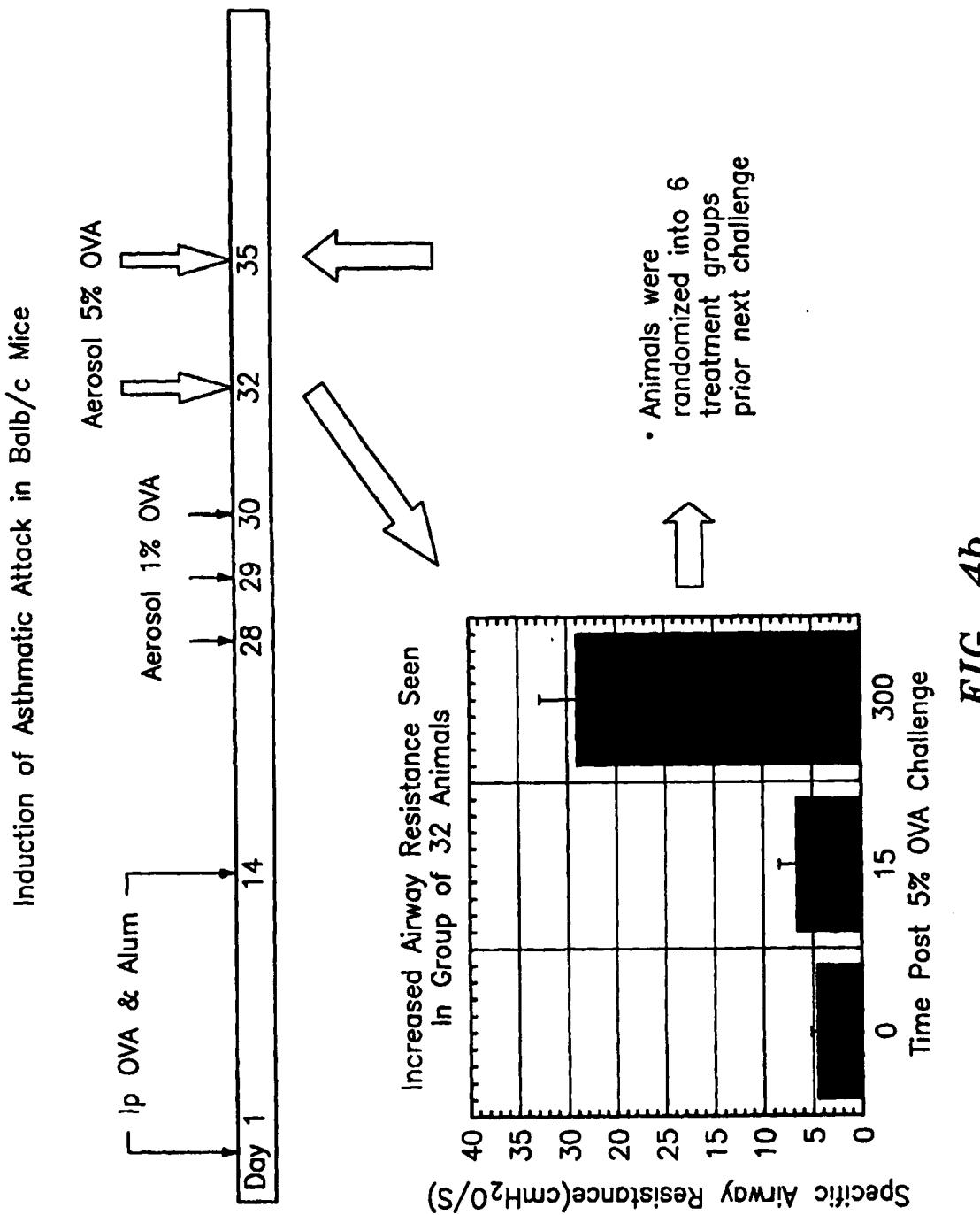


FIG. 4b

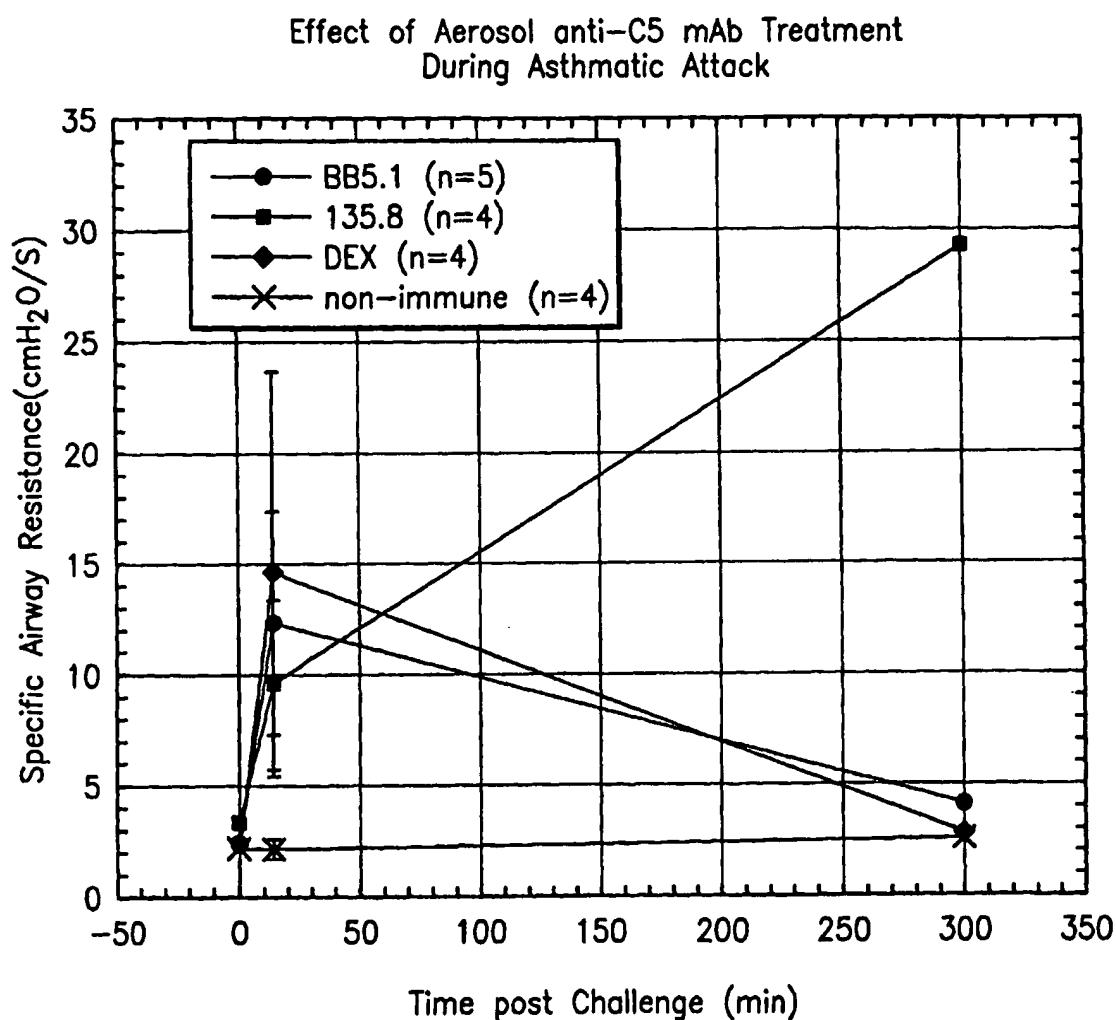


FIG. 5a

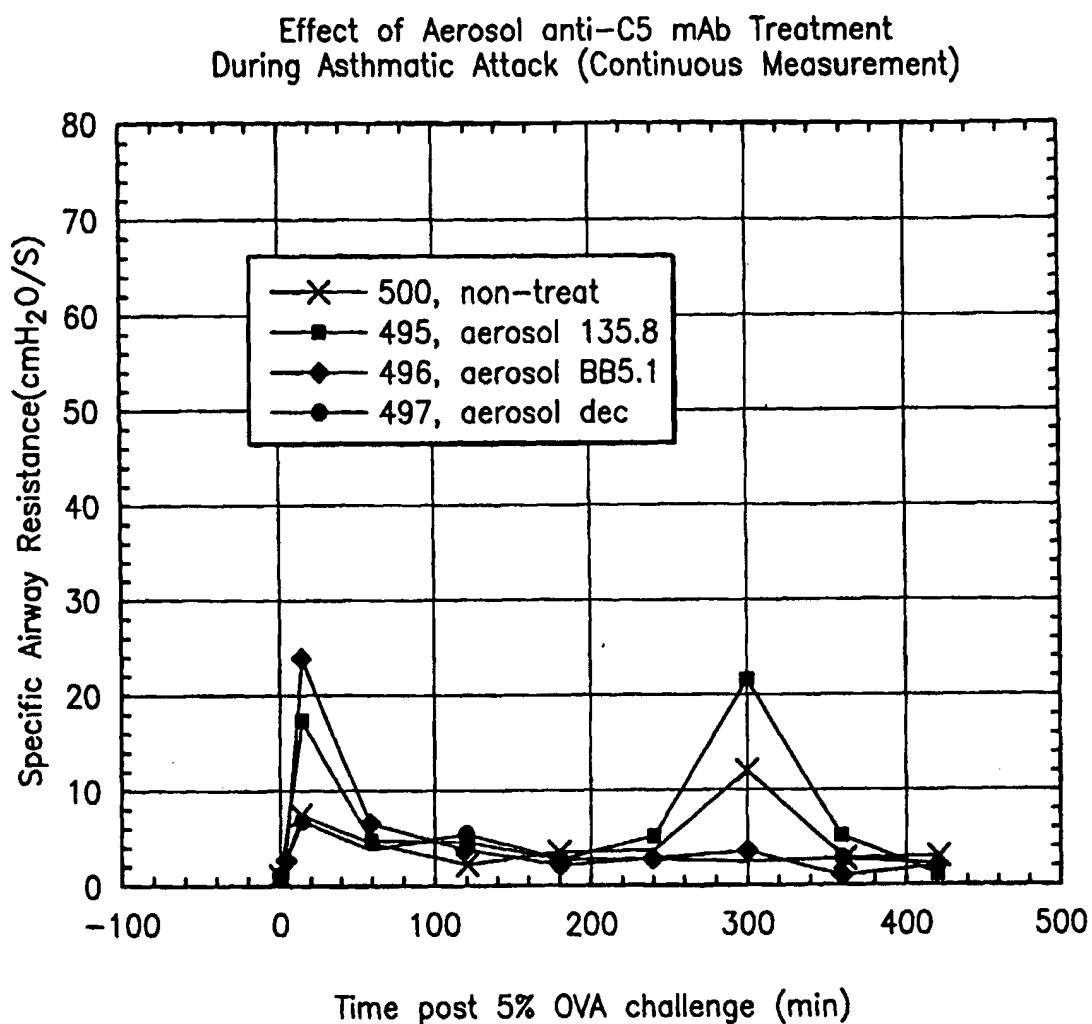


FIG. 5b

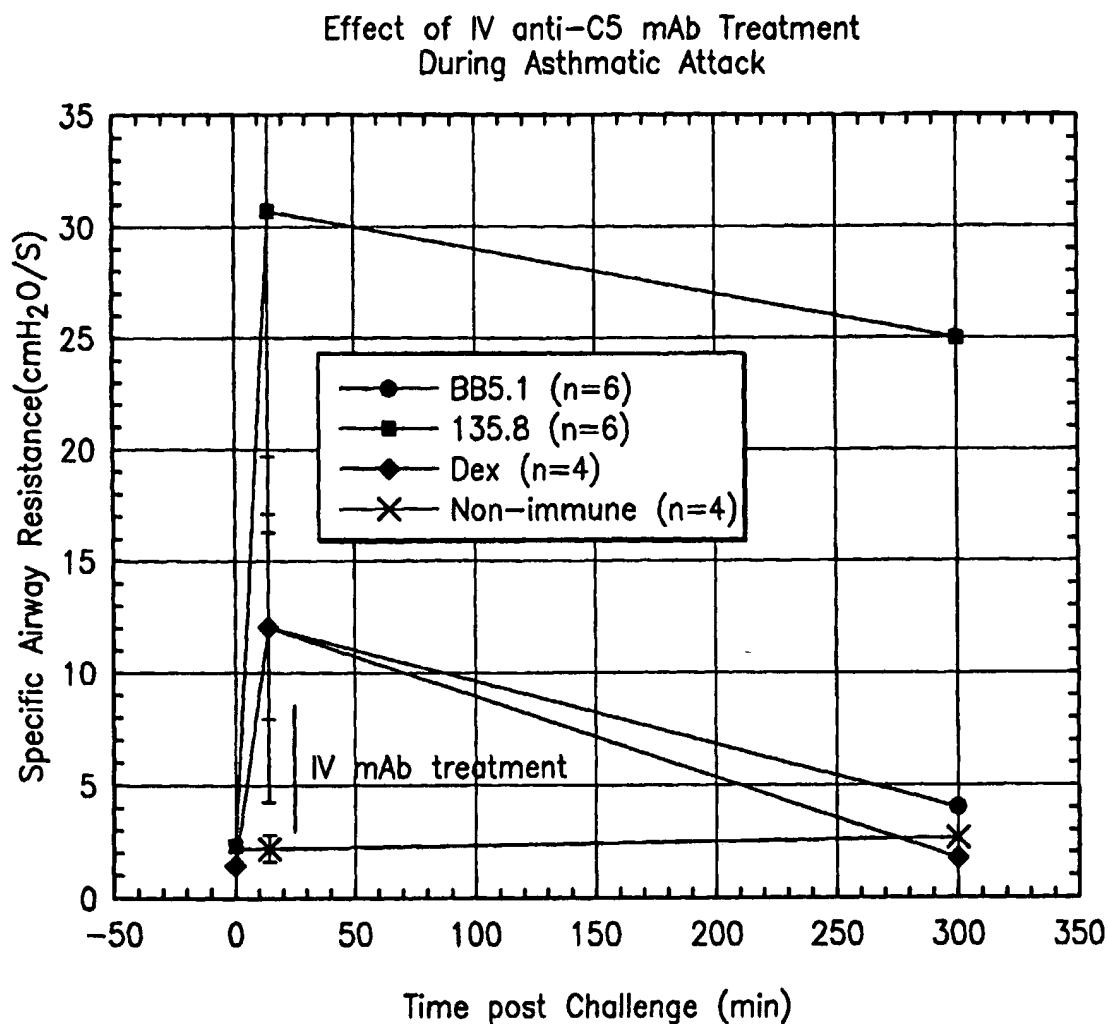


FIG. 6a

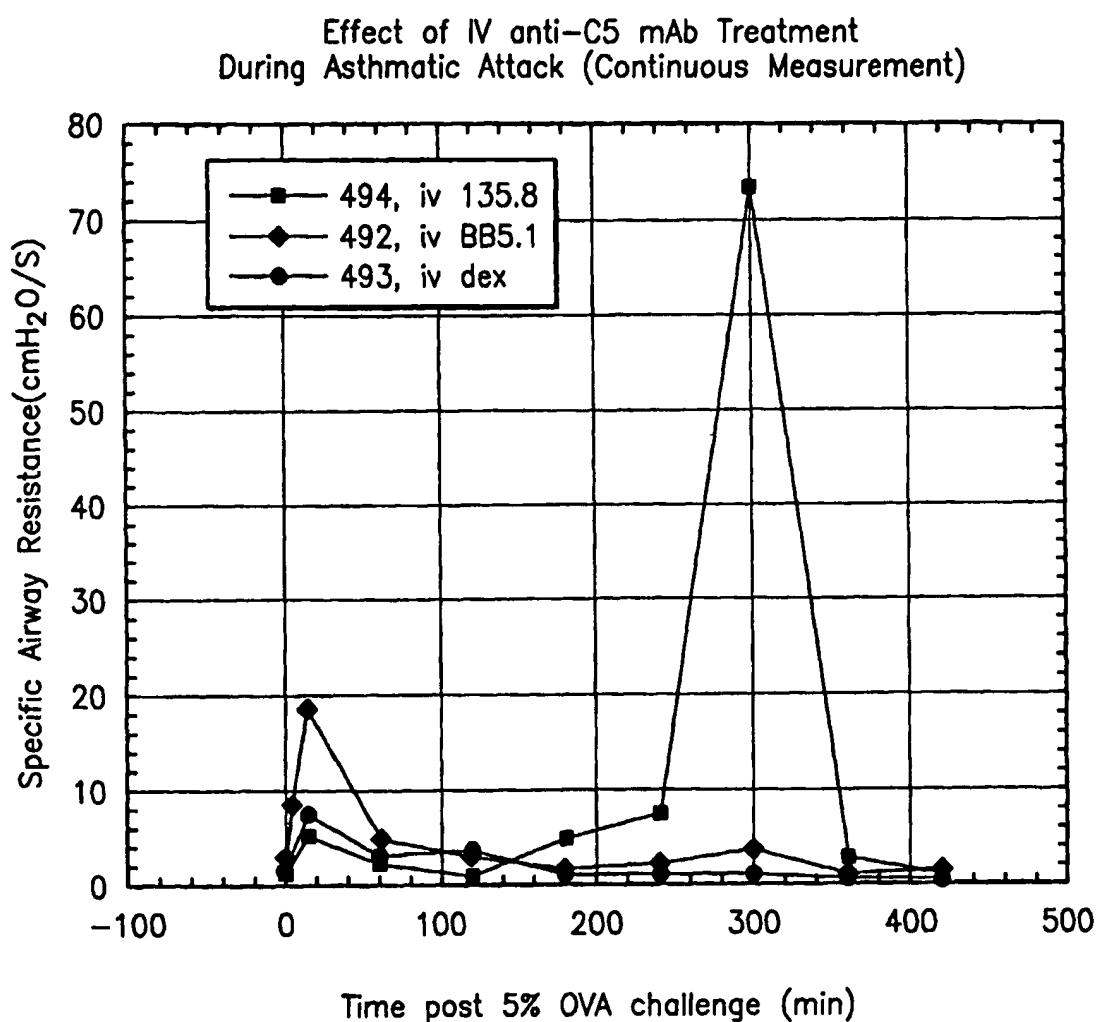


FIG. 6b

11/19

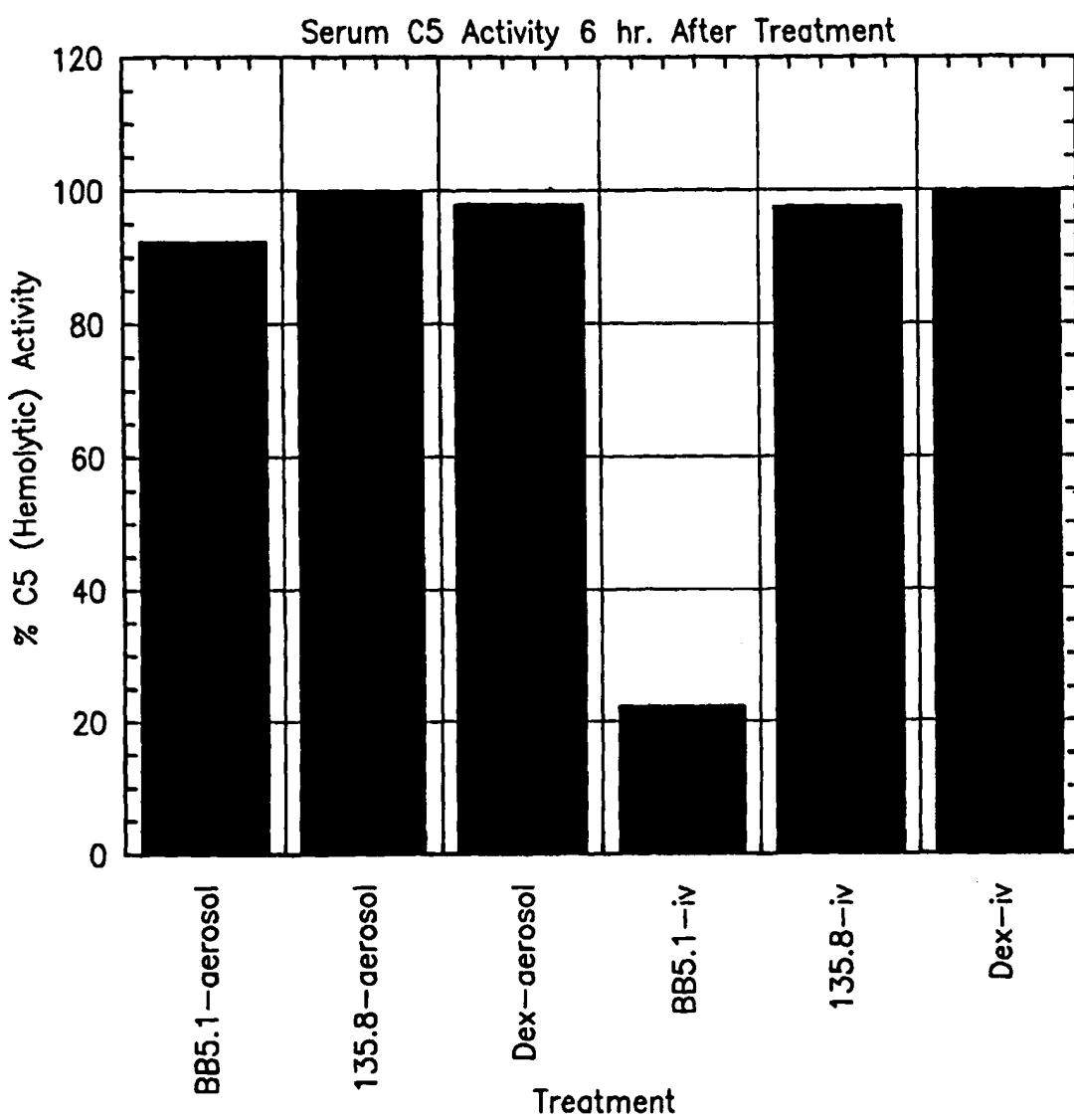


FIG. 7

12/19

Effect of Steroid and anti-C5 Treatment on BAL total WBC Count
(5hr after OVA challenge and mAb treatment)

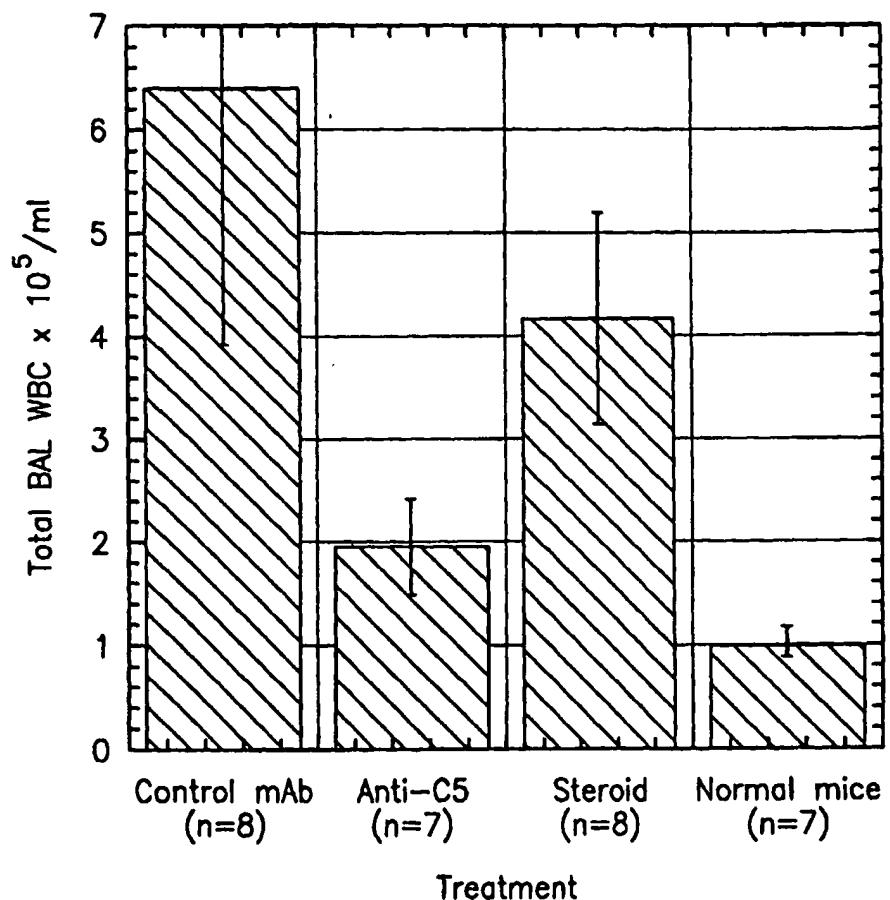
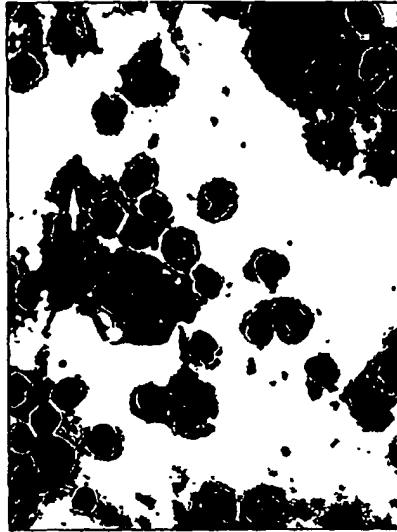


FIG. 8

Eosinophils are the predominant Inflammatory cells in BAL



Non-immunized
FIG. 9a



Control mAb
FIG. 9b



Steroid
FIG. 9c



Anti-C5
FIG. 9d

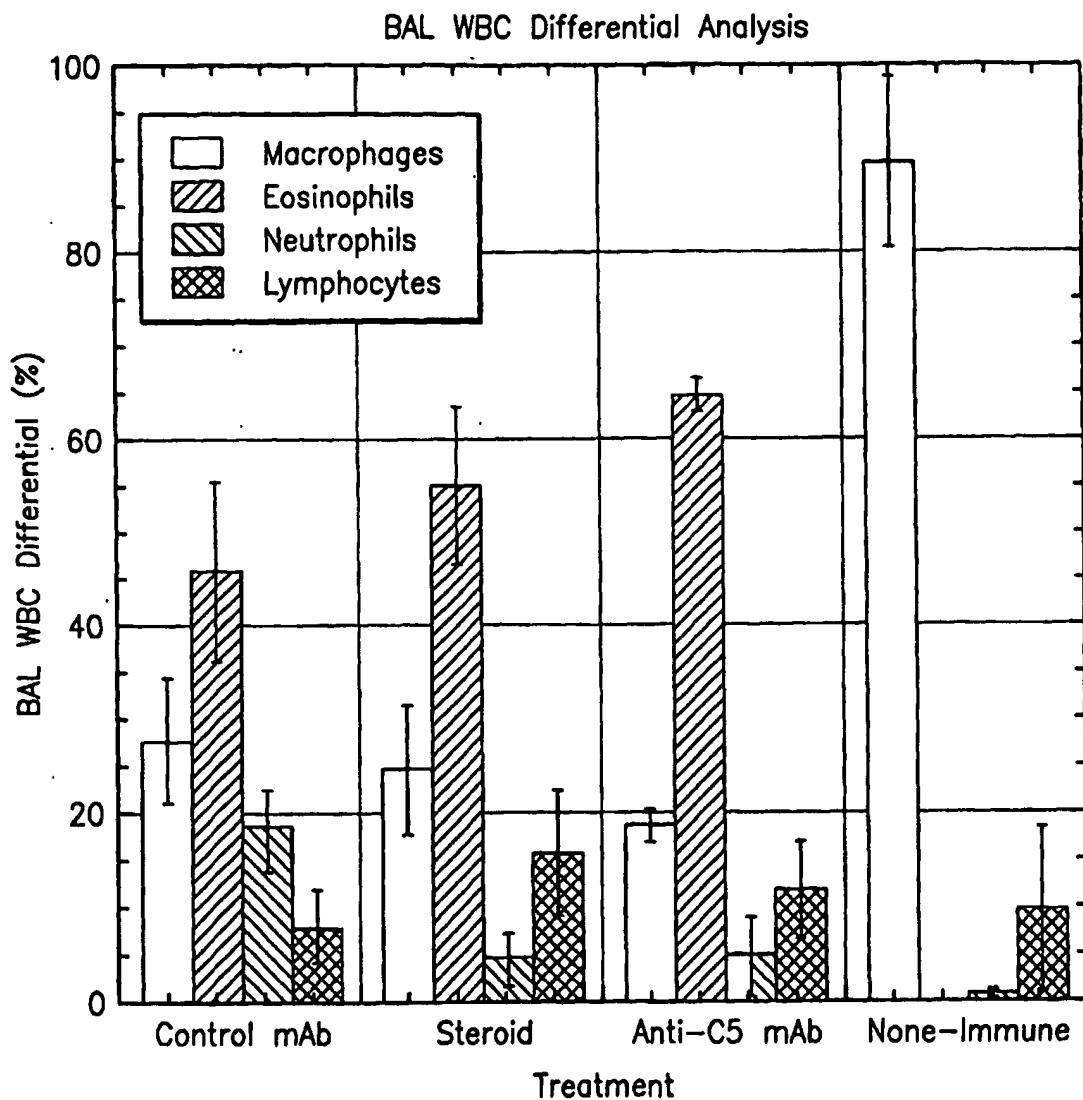


FIG. 10

Effect of Steroid and anti-C5 Treatment on BAL Histamine Level
(5hr after OVA challenge and mAb treatment)

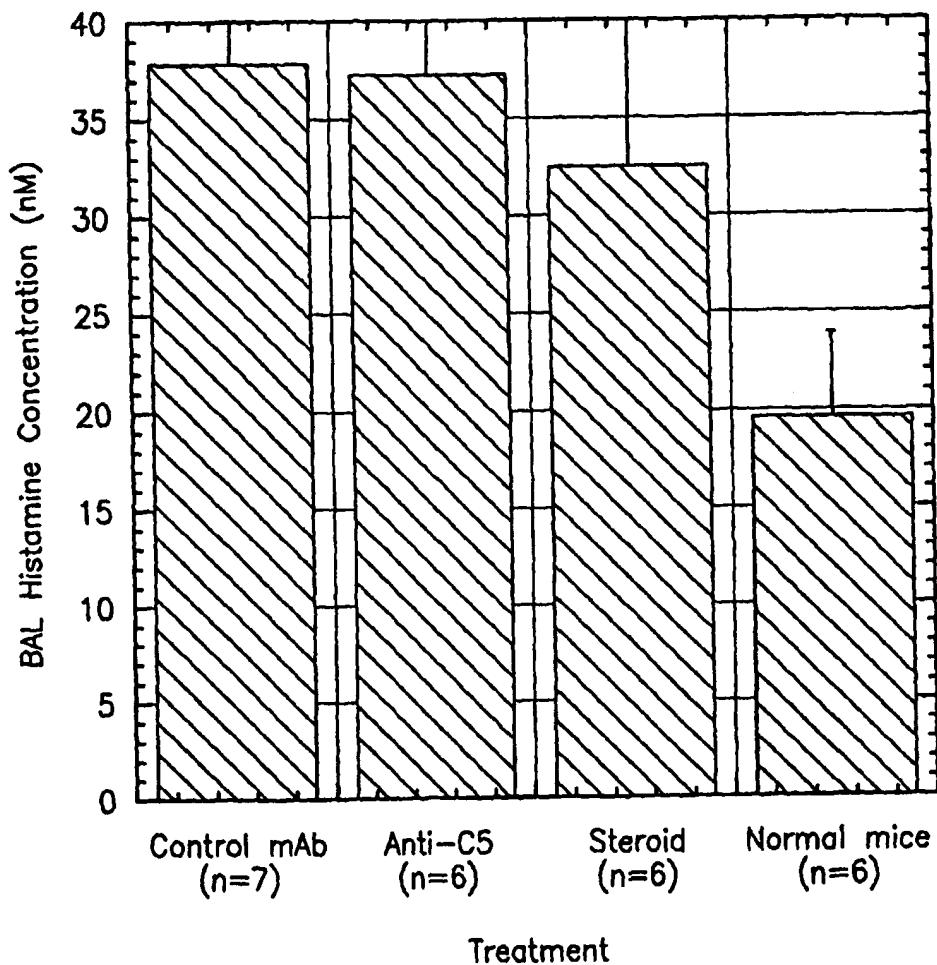


FIG. 11

New Data: Effect of Steroid and anti-C5 Treatment on BAL MMP-9 Level
(5hr after OVA challenge and mAb treatment)

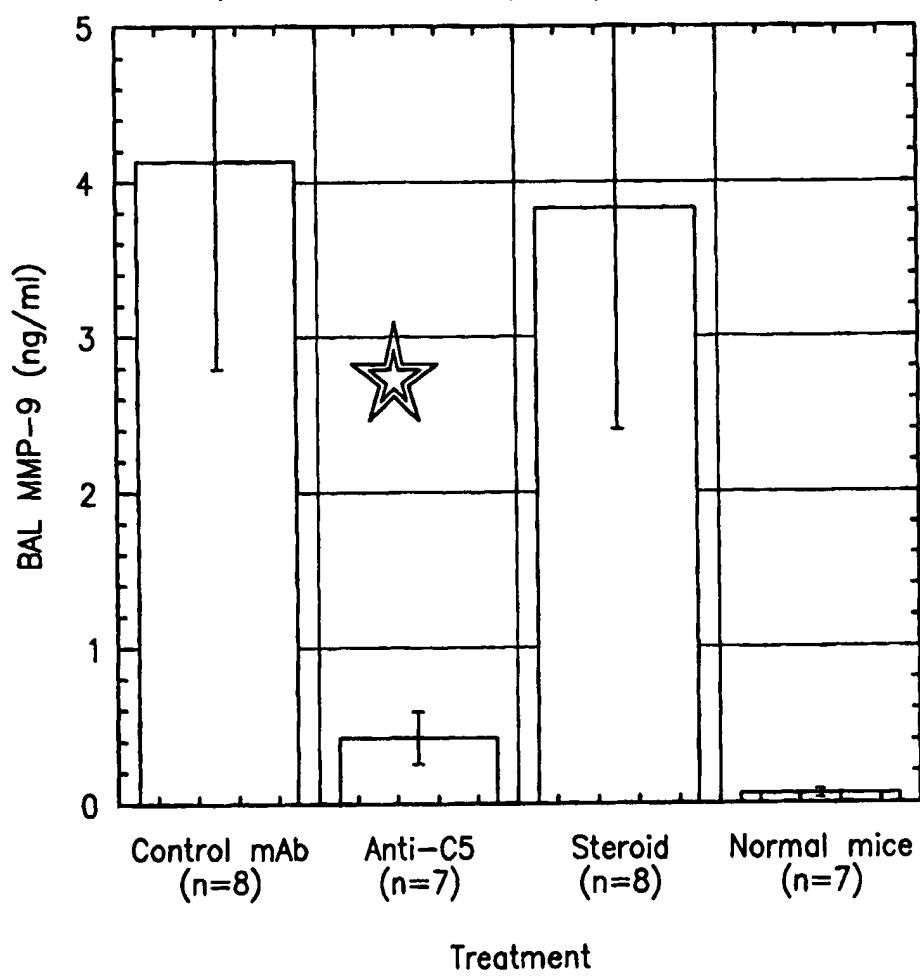


FIG. 12

17/19

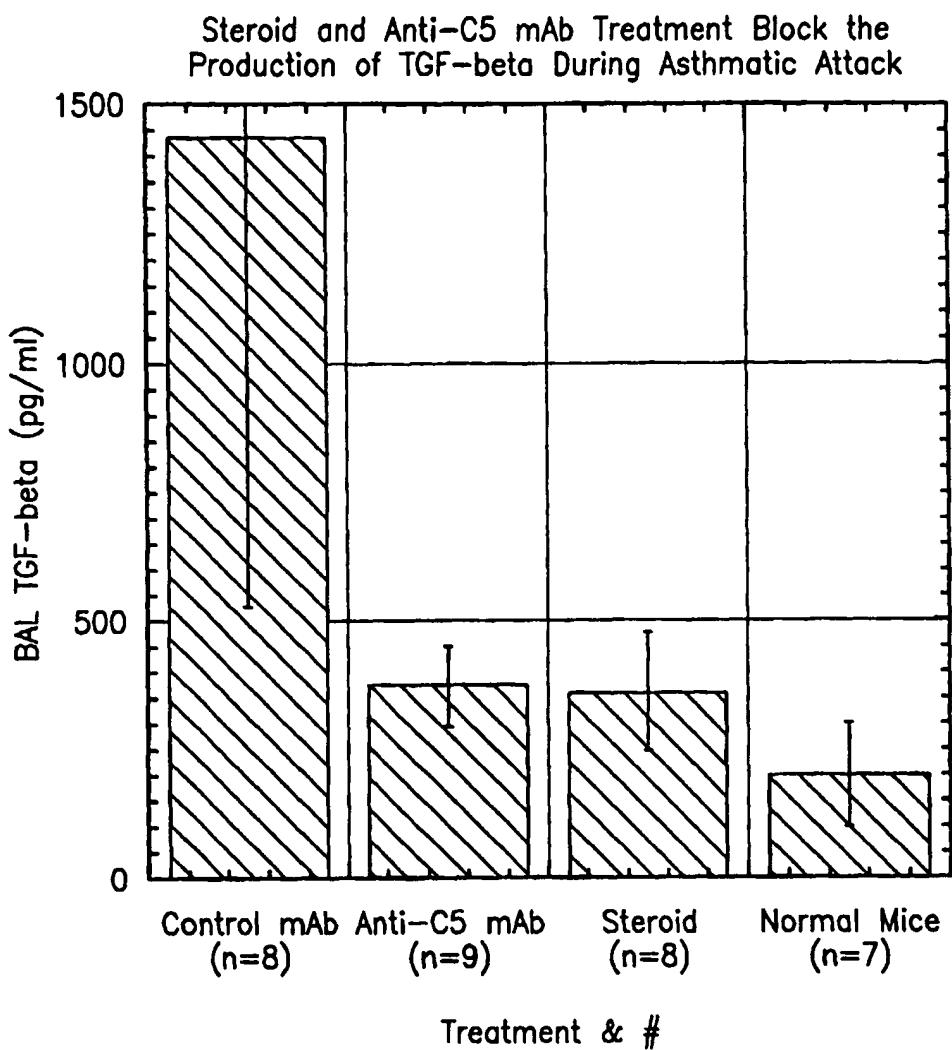


FIG. 13

18/19

Test the Direct & Immediate Bronchial Dilation Effect of C5 Inhibitor Or In Combination with Salbutamol During Wheezing (Provocation with Allergen)

Cannulation & Aerosol C5 inhibitor or β_2 agonist & Testing RL simultaneously

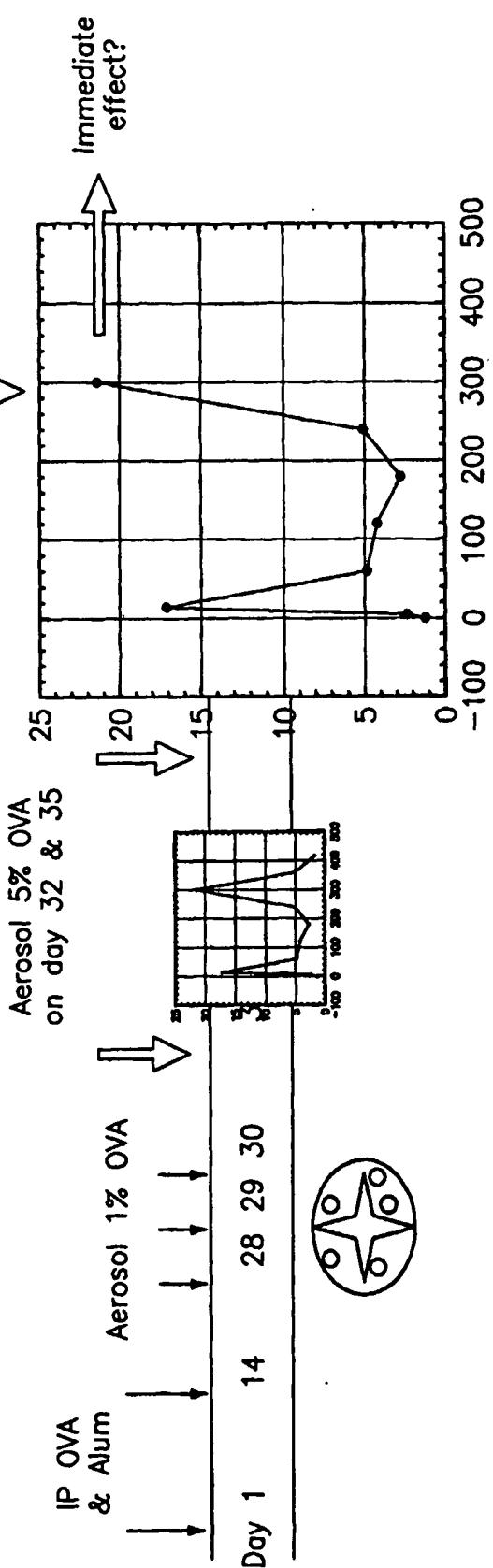


FIG. 14

Test the Direct & Immediate Bronchial Dilation Effect of C5 Inhibitor Or In Combination with Salbutamol During Wheezing

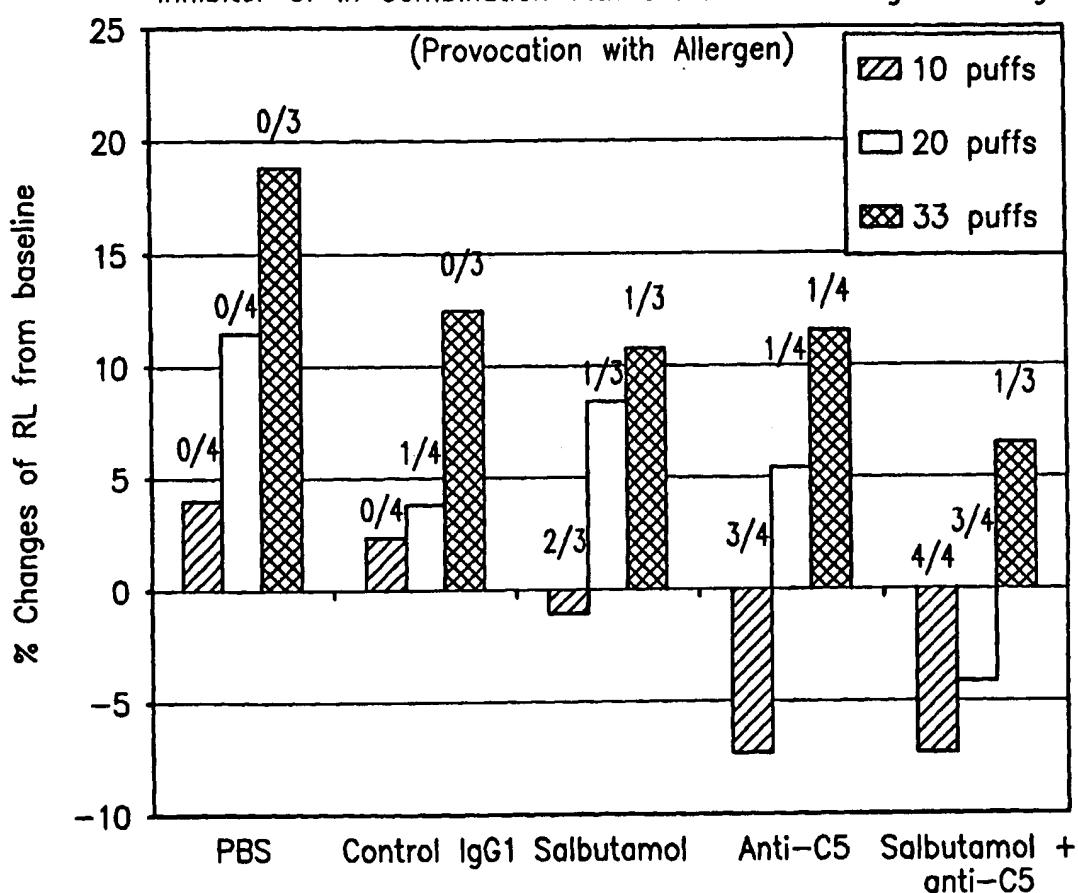


FIG. 15

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US03/27808

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : A61K 39/395; C07K 16/36
 US CL : 424/130.1, 133.1, 141.1, 145.1, 158.1; 530/387.1, 387.3, 388.1, 388.25

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
 U.S. : 424/130.1, 133.1, 141.1, 145.1, 158.1; 530/387.1, 387.3, 388.1, 388.25

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
 NONE

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
 Please See Continuation Sheet

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 6,355,245 B1 (EVANS et al.) 12 March 2002 (12 March 2002), see entire document.	1-44
Y	US 6,316,502 B1 (LAI et al.) 13 November 2001 (13.11.2001), see entire document, including column 15, lines 34-39 and column 15, line 66 - column 16, line 2).	1-44
Y	EP 0 649 468 B1 (SMITH et al.) 15 March 2000 (15.03.2000), see entire document, including page 6, line 22.	1-44

Further documents are listed in the continuation of Box C.

See patent family annex.

• Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier application or patent published on or after the international filing date	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&"	document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means		
"P" document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

10 December 2003 (10.12.2003)

Date of mailing of the international search report

12 FEB 2004

Name and mailing address of the ISA/US

Mail Stop PCT, Attn: ISA/US
 Commissioner for Patents
 P.O. Box 1450
 Alexandria, Virginia 22313-1450
 Facsimile No. (703)305-3230

Authorized officer

Phillip Gambel

Telephone No. 703-308-0196

INTERNATIONAL SEARCH REPORT

PCT/US03/27808

Continuation of B. FIELDS SEARCHED Item 3:
DIALOG, BIOSIS, CA, EMBASE, MEDLINE, WEST
search terms: c5, c5a, antibod?, 5g1.1, asthma, bronchial, pulmonary, wang